Listing of Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application.

What is claimed is:

- 1 1. (Original) A partially thio-modified aptamer that binds to a TGF-beta protein.
- 1 2. (Original) The aptamer of claim 1, wherein the TGF-beta protein comprises a
- 2 human TGF-beta.
- 1 3. (Original) The aptamer of claim 1, wherein the TGF-beta protein comprises a
- 2 TGF-beta dimer.
- 1 4. (Original) The aptamer of claim 3, wherein the TGF-beta dimer is a homodimer.
- 1 5. (Original) The aptamer of claim 4, wherein the TGF-beta homodimer is a TGF-
- 2 beta 1, 2 or 3 homodimer.
- 1 6. (Original) The aptamer of claim 3, wherein the TGF-beta dimer is a TGFbeta 1, 2
- 2 or 3 heterodimer.
- 1 7. (Currently Amended) The aptamer of claim 1, wherein the aptamer comprises one
- 2 or more thio-modifications as set forth in the sequence and modifications of SEQ ID
- 3 NO[[S]]: 62.
- 1 8. (Original) The aptamer of claim 1, wherein the aptamer is achiral.
- 1 9. (Original) The aptamer of claim 1, wherein the aptamer further comprises a
- detectable label.
- 1 10. (Original) The aptamer of claim 1, further comprising one or more
- 2 pharmaceutically acceptable salts.
- 1 11. (Original) The aptamer of claim 1, further comprising a diluent.

- 1 12. (Withdrawn) A partially thio-modified aptamer that binds to a TGF-beta receptor.
- 1 13. (Withdrawn) The aptamer of claim 12, wherein the TGF-beta receptor is a
- 2 signaling receptor.
- 1 14. (Withdrawn) The aptamer of claim 12, wherein the TGF-beta receptor is a co-
- 2 receptor.
- 1 15. (Withdrawn) The aptamer of claim 13, wherein the TGF-beta signaling receptor
- 2 comprises a human TGF-beta signaling receptor.
- 1 16. (Withdrawn) The aptamer of claim 13 wherein the TGF-beta signaling receptor
- 2 comprises a TbetaRI or a TbetaRII receptor.
- 1 17. (Withdrawn) The aptamer of claim 13, wherein the target of the aptamer is the GS
- 2 domain of a TbetaRI receptor.
- 1 18. (Withdrawn) The aptamer of claim 14, where the co-receptor is TGF-beta 3.
- 1 19. (Withdrawn) The aptamer of claim 12, wherein the aptamer is achiral.
- 1 20. (Withdrawn) A partially thio-modified aptamer that binds to a ligand-receptor
- 2 complex comprising a TGF-beta ligand and a receptor complex comprising a TbetaRI
- 3 and a TbetaRII receptors.
- 1 21. (Withdrawn) The aptamer of claim 20, wherein the target of the aptamer is the GS
- 2 domain of a TbetaRI receptor.
- 1 22. (Withdrawn) The aptamer of claim 20, wherein the aptamer is achiral.
- 1 23. (Withdrawn) A partially thio-modified aptamer that binds to a ligand binding trap
- 2 capable of trapping TGF-beta ligands.
- 1 24. (Withdrawn) The aptamer of claim 23, wherein the ligand binding trap comprises
- decorin, latency-associated protein (LAP) or alpha-macroglobulin.

- 1 25. (Withdrawn) The aptamer of claim 23, wherein the aptamer is achiral.
- 1 26. (Withdrawn) A partially thio-modified aptamer that binds to an auxiliary protein
- 2 that promotes binding of TGF-beta ligand to Tbeta signaling receptors.
- 1 27. (Withdrawn) The aptamer of claim 26, wherein the auxiliary protein is a SARA
- 2 protein.
- 1 28. (Withdrawn) The aptamer of claim 26, wherein the aptamer is achiral.
- 1 29. (Withdrawn) A partially thio-modified aptamer that binds to a Smad protein.
- 1 30. (Withdrawn) The aptamer of claim 29, wherein the Smad protein is an R-Smad, a
- 2 Co-Smad, an I-Smad or a combination thereof.
- 1 31. (Withdrawn) The aptamer of claim 29, wherein the aptamer is achiral.
- 1 32. (Withdrawn) A partially thio-modified aptamer that binds to a TGF-beta protein
- 2 complex and enhances TGF-beta activity.
- 1 33. (Withdrawn) The aptamer of claim 32, wherein the binding site of the aptamer on
- 2 the TGF-beta protein complex comprises a region of a ligand binding trap protein.
- 1 34. (Withdrawn) The aptamer of claim 32, wherein the binding site of the aptamer on
- 2 the TGF-beta protein complex comprises a region of an inhibitory I-Smad.
- 1 35. (Withdrawn) The aptamer of claim 32, wherein the aptamer is achiral.
- 1 36. (Withdrawn) A partially thio-modified aptamer that binds to a TGF-beta protein
- 2 complex and inhibits TGF-beta activity.
- 1 37. (Withdrawn) The aptamer of claim 36, wherein the binding site of the aptamer on
- 2 the TGF-beta protein complex comprises a region of an R-Smad or a Co-Smad.
- 1 38. (Withdrawn) The aptamer of claim 36, wherein the aptamer is achiral.

- 1 39. (Withdrawn) A partially modified thioaptamer that inhibits TGF-beta activity by
- 2 binding to a TGF-beta ligand, a TGF-beta ligand-Tbeta receptor complex, a TGF-beta
- 3 signaling receptor and co-receptor, to an R-Smad or a Co-Smad.
- 1 40. (Withdrawn) The aptamer of claim 39, wherein the aptamer is achiral.
- 1 41. (Withdrawn) A partially modified thioaptamer that modifies TGF-beta activity by
- 2 binding to a TGF-beta ligand, a TGF-beta ligand-Tbeta receptor complex, a TGF-beta
- 3 signaling receptor and co-receptor, to an R-Smad or a Co-Smad.
- 1 42. (Withdrawn) A method of inhibiting TGF-β activity comprising the steps of:
- 2 providing to a host in need of therapy a pharmaceutically effective amount of a
- 3 thioaptamer that specifically binds to and inhibits TGF-β activity.
- 1 43. (Withdrawn) The method of claim 42, wherein the thioaptamer is provided to the
- 2 host to ameliorate the effects of: fibrosis, scarring and adhesion during wound healing;
- 3 fibrotic diseases of the lung, liver and kidney; atherosclerosis, arteriosclerosis; cancers
- 4 including gliomas, colon cancer, prostate cancer, breast cancer, neurofibromas, lung
- 5 cancer; angiopathy, vasculopathy, nephropathy; systemic sclerosis; viral infections
- 6 accompanied by immune suppression (HIV, HCV); and immunological disorders and
- 7 deficiencies (auto-immune diseases).
- 1 44. (Withdrawn) A method of quantitating TGF-β levels in a sample comprising the
- 2 step of contacting a sample with a TGF-β-specific thioaptamer.
- 1 45. (Withdrawn) The method of claim 44, wherein the samples comprises a
- 2 physiological sample.
- 1 46. (Withdrawn) The method of claim 44, wherein the sample comprise a blood,
- 2 tissue, cells, supernatant, media.

- 1 47. (Withdrawn) The method of claim 44, wherein the TGF-β protein comprises a
- 2 human TGF-β.
- 1 48. (Withdrawn) The method of claim 44, wherein the TGF-β protein comprises a
- 2 TGF-β homodimer.
- 1 49. (Withdrawn) The method of claim 44, wherein the TGF-β protein comprises a
- 2 TGF- β 1, 2 or 3 heterodimer.
- 1 50. (Withdrawn) The method of claim 44, wherein the thioaptamer comprises one or
- 2 more thio-modifications as set forth in SEQ ID NOS.: 4-22.
- 1 51. (Withdrawn) The method of claim 44, wherein the thioaptamer further comprises
- 2 a detectable label.
- 1 52. (Withdrawn) The method of claim 44, wherein the thioaptamer further comprises
- 2 a detectable detectable selected from the group consisting of a colorimetric, a fluorescent,
- 3 a radioactive and an enzymatic agent.
- 1 53. (Withdrawn) A method of modulating TGF-β signaling comprising the steps of:
- 2 administering to a host a TGF- β specific thioaptamer that modulates the activity through
- 3 the TGF- β receptor in a dosage effective to reduce activity of the TGF- β .
- 1 54. (Withdrawn) The method of claim 53, wherein the thioaptamer modulates the
- 2 activity through the TGF- β receptor by increasing activity.
- 1 55. (Withdrawn) The method of claim 53, wherein the thioaptamer modulates the
- 2 activity through the TGF-β receptor by decreasing activity.
- 1 56. (Withdrawn) The method of claim 53, wherein the thioaptamer is selected from
- 2 the group consisting of SEQ ID NOS.:4-22.

- 1 57. (Withdrawn) A method of treating a pathological condition due to increased TGF-
- 2 β activity comprising the steps of:
- 3 administering to a host an effective dosage of a thioaptamer that modulates TGF-β.
- 1 58. (Withdrawn) The method of claim 57, wherein the thioaptamer binds to TGF-β,
- 2 the TGF-β receptor, a TGF-β auxiliary protein, a TGF-β ligand binding trap protein or a
- 3 TGF-β Smad protein.
- 1 59. (Withdrawn) The method of claim 57, wherein the thioaptamer modulates the
- 2 activity through the TGF- β receptor by increasing activity.
- 1 60. (Withdrawn) The method of claim 57, wherein the thioaptamer modulates the
- 2 activity through the TGF-β receptor by decreasing activity.
- 1 61. (Withdrawn) The method of claim 57, wherein the thioaptamer is selected from
- 2 the group consisting of SEQ ID NOS.: 4-22.
- 1 62. (Withdrawn) The method of claim 57, wherein the pathological condition
- 2 comprises:
- 3 fibrosis, scarring and adhesion during wound healing; fibrotic diseases of the lung, liver
- 4 and kidney; atherosclerosis and arteriosclerosis; cancers such as gliomas, colon cancer,
- 5 prostate cancer, breast cancer, neurofibromas, lung cancer; angiopathy, vasculopathy,
- 6 nephropathy; systemic sclerosis; viral infections accompanied by immune suppression
- 7 (HIV, HCV); and immunological disorders and deficiencies (auto-immune diseases).
- 1 63. (Withdrawn) The method of claim 57, wherein the TGF-β specific thioaptamer is
- 2 encapsulated.
- 1 64. (Withdrawn) The method of claim 57, wherein the capsule is degradable by an
- 2 external stimulus to release the TGF- β specific thioaptamer.

- 1 65. (Withdrawn) The method of claim 57, wherein the external stimulus is selected
- 2 from the group consisting of UV light, acid, water, in vivo enzymes, ultrasound and heat.
- 1 66. (Withdrawn) The method of claim 57, wherein the TGF-β specific thioaptamer is
- 2 bound to a binding molecule.
- 1 67. (Withdrawn) The method of claim 57, wherein the TGF-β specific thioaptamer is
- 2 bound to a binding molecule and further comprising the step of detaching the binding
- 3 molecule from the TGF- β specific thioaptamer.
- 1 68. (Withdrawn) A method of treating a pathological condition in which increased
- 2 TGF-β activity has been implicated comprising the steps of:
- 3 administering to a host a TGF-β specific thioaptamer in a pharmaceutically acceptable
- 4 carrier at a dosage effective to reduce TGF-β activity.
- 1 69. (Withdrawn) The method of claim 68, wherein the pharmaceutically acceptable
- 2 carrier is selected from the group consisting of a cream, gel, aerosol and powder for
- 3 topical application.
- 1 70. (Withdrawn) The method of claim 68, wherein the pharmaceutically acceptable
- 2 carrier is selected from the group consisting of a sterile solution for injection, irrigation
- 3 and inhalation.
- 1 71. (Withdrawn) The method of claim 68, wherein the pharmaceutically acceptable
- 2 carrier comprises a sterile dressing for topically covering a wound.
- 1 72. (Withdrawn) The method of claim 68, wherein the pharmaceutically acceptable
- 2 carrier is selected from the group consisting of a biopolymer and a polymer for
- 3 implanting within a wound.
- 1 73. (Withdrawn) The method of claim 68, further comprising the step of
- 2 administering a growth factor other than TGF-β.

- 1 74. (Withdrawn) The method of claim 68, wherein the TGF-β specific thioaptamer is
- 2 encapsulated.
- 1 75. (Withdrawn) A method of modulating TGF-β signaling comprising the steps of:
- 2 administering to a host a TGF-β ligand binding trap specific thioaptamer that modulates
- 3 the activity through the TGF- β receptor in a dosage effective to reduce activity of the
- 4 TGF-β.
- 1 76. (Withdrawn) A method of modulating TGF-β signaling comprising the steps of:
- 2 administering to a host a TGF-β auxiliary protein specific thioaptamer that modulates the
- 3 activity through the TGF- β receptor in a dosage effective to reduce activity of the TGF- β .
- 1 77. (Withdrawn) A method of modulating TGF-β signaling comprising the steps of:
- 2 administering to a host a TGF-β Smad protein specific thioaptamer that modulates the
- activity through the TGF- β receptor in a dosage effective to reduce activity of the TGF- β .
- 1 78. (NEW) A partially thio-modified aptamer that binds specifically to TGF-β
- 2 comprising a sequence and modifications that is at least 80% complementary to SEQ ID
- 3 NO: 62.